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	ENTRY	SESSION
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	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	78.51	97.82

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-5.10	-5.10

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FILE COVERS 1907 - 10 Sep 2010 VOL 153 ISS 12
FILE LAST UPDATED: 9 Sep 2010 (20100909/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2010
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2010

HCAplus now includes complete International Patent Classification (IPC) reclassification data for the third quarter of 2010.

Updated Search

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<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s sandmeyer () reaction? and copper?

1828 SANDMEYER

3 SANDMEYERS

1829 SANDMEYER

(SANDMEYER OR SANDMEYERS)

4722051 REACTION?

1249 SANDMEYER (W) REACTION?

1123055 COPPER?

L11 97 SANDMEYER (W) REACTION? AND COPPER?

=> s copper () II

1120963 COPPER

544 COPPERS

1121038 COPPER

(COPPER OR COPPERS)

2383058 II

1255 IIS

2383799 II

(II OR IIS)

L12 52536 COPPER (W) II

=> s l12 and sandmeyer () reaction?

1828 SANDMEYER

3 SANDMEYERS

1829 SANDMEYER

(SANDMEYER OR SANDMEYERS)

4722051 REACTION?

1249 SANDMEYER (W) REACTION?

L13 9 L12 AND SANDMEYER (W) REACTION?

=> s l13 and review/dt

2416647 REVIEW/DT

L14 0 L13 AND REVIEW/DT

=> d l13, ibib abs, 1-9

THE ESTIMATED COST FOR THIS REQUEST IS 27.90 U.S. DOLLARS

DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N:y

L13 ANSWER 1 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2009:1149701 HCAPLUS

DOCUMENT NUMBER: 151:448008

TITLE: Chelation-Assisted Palladium-Catalyzed Direct
Cyanation of 2-Arylpyridine C-H Bonds

AUTHOR(S): Jia, Xiaofei; Yang, Dongpeng; Zhang, Shouhui; Cheng,
Jiang

CORPORATE SOURCE: College of Chemistry and Materials Engineering,
Wenzhou University, Wenzhou, 325027, Peop. Rep. China

SOURCE: Organic Letters (2009), 11(20), 4716-4719

CODEN: ORLEF7; ISSN: 1523-7060

Updated Search

STNsdddaeatat

PUBLISHER: American Chemical Society
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 151:448008

AB A chelation-assisted palladium-catalyzed ortho-cyanation of the sp² C-H bond of arylpyridine derivs. by CuCN provided aromatic nitriles in moderate to good yields. Notably, the reaction could be conducted on a 10 mmol scale. The key intermediate of a natural product of *Menispermum dauricum* DC was concisely synthesized by the described procedure. This new approach represents an exceedingly practical method for the synthesis of aromatic nitriles and offers an attractive alternative to the traditional Sandmeyer reaction.

OS.CITING REF COUNT: 12 THERE ARE 12 CAPLUS RECORDS THAT CITE THIS RECORD (12 CITINGS)

REFERENCE COUNT: 77 THERE ARE 77 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 2 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2008:383494 HCAPLUS

DOCUMENT NUMBER: 148:552026

TITLE: Synthesis of dioxin-like monofluorinated PCBs: for the use as internal standards for PCB analysis

AUTHOR(S): Sott, Richard; Hawner, Christine; Johansen, Jon E.

CORPORATE SOURCE: Chiron AS, Trondheim, N-7041, Norway

SOURCE: Tetrahedron (2008), 64(18), 4135-4142

CODEN: TETRAB; ISSN: 0040-4020

PUBLISHER: Elsevier Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 148:552026

AB Monofluorinated polychlorinated biphenyls (fluoro-PCBs) were prepared using the Suzuki-coupling, for use as anal. stds. for PCB measurements. Seven of these fluoro-PCBs are analogs of the dioxin-like PCBs, listed by the WHO as the most toxic PCB congeners. Four highly chlorinated fluoro-PCBs were prepared by Suzuki-coupling of 2,3,5,6-tetrachloro-bromoaniline with various substituted arylboronic acids. The resulting amino-fluoro-PCBs are chlorinated using the Sandmeyer reaction or deaminated to yield tetra-, penta- and hexa-chlorinated fluoro-PCBs. The fluoro-PCBs elute just before the corresponding PCBs in the GC chromatogram, which strongly indicates their potential as anal. stds.

OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 3 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2004:625843 HCAPLUS

DOCUMENT NUMBER: 141:157033

TITLE: Process for the preparation of iodochromones, intermediates in the synthesis of fungicides, by condensation of nitriles with o-hydroxyaryl esters, cyclization, diazotization, decomposition of the diazonium salt, etherification and/or transesterification

INVENTOR(S): Delamare, Madelaine; Casado, Michel

PATENT ASSIGNEE(S): Bayer Cropscience S. A., Fr.

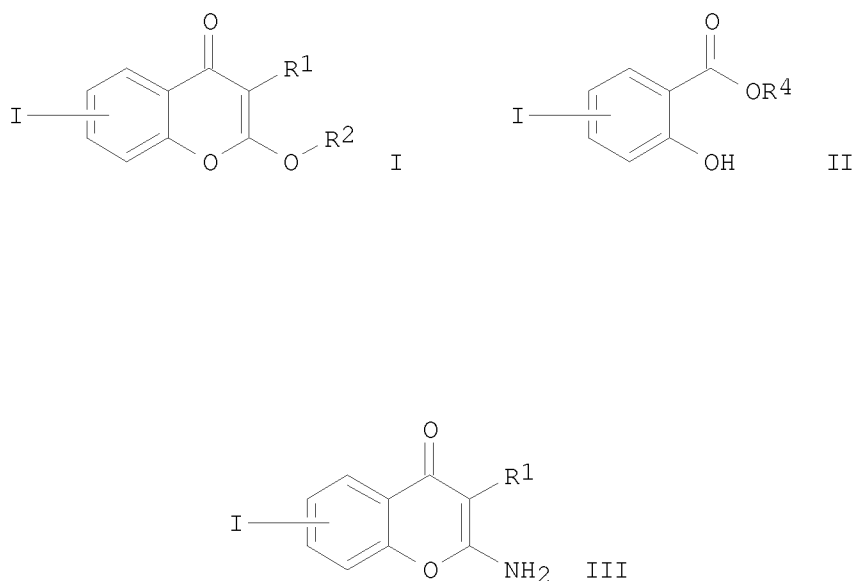
Updated Search

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SOURCE: Eur. Pat. Appl., 16 pp.
CODEN: EPXXDW
DOCUMENT TYPE: Patent
LANGUAGE: French
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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EP 1443044	A1	20040804	EP 2003-356014	20030203
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
WO 2004069821	A1	20040819	WO 2004-EP1786	20040128
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI				
RW: BW, GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1592681	A1	20051109	EP 2004-705816	20040128
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
BR 2004006687	A	20051220	BR 2004-6687	20040128
CN 1745075	A	20060308	CN 2004-80003356	20040128
JP 2006516595	T	20060706	JP 2006-501930	20040128
IN 2005DN03345	A	20070420	IN 2005-DN3345	20050727
MX 2005008277	A	20050920	MX 2005-8277	20050803
US 20060111434	A1	20060525	US 2005-544294	20051017
PRIORITY APPLN. INFO.:			EP 2003-356014	A 20030203
			WO 2004-EP1786	W 20040128
ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT				
OTHER SOURCE(S):			CASREACT 141:157033; MARPAT 141:157033	
GI				

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AB The invention is related to a process for the preparation of iodochromones (I), intermediates in the synthesis of fungicides, by condensation of nitriles R^1CH_2CN with o-hydroxyaryl esters (II), cyclization in aqueous acidic media, diazotization of the amine (III) with alkyl(thio)nitrites, or alkylthionitrates, or $NaNO_2$ in HCl, or H_2SO_4 , Sandmeyer reaction of the in-situ formed diazonium salt in the presence of copper halides or $Cu(OR_3)$, followed by etherification with or transesterification with sodium alkoxides [R^1 , R^2 , R^3 , R^4 = independently (un)substituted alk(en/yn)yl, carbocyclyl, heterocyclyl]. The advantages include high reaction yields, and conversions, use of cheap reagents, reduction in number of steps, and elimination of undesired steps and byproduct formation. For example, 2-butoxy-6-iodo-3-propylchromone (m.p. = 69-71°) was prepared by condensation of valeronitrile with 4-iodosalicylate in the presence of DIPA in THF/cyclization in the presence of NH_4Cl , diazotization/Sandmeyer reaction in the presence of t-Bu nitrite/ $CuCl_2/CH_3CN$, and etherification of the chlorochromone with sodium butylate in n-butanol.

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 4 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 2002:153725 HCAPLUS

DOCUMENT NUMBER: 137:124967

TITLE: Synthesis of fluorine containing polyfunctional aromatic compounds. Sandmeyer synthesis of 4-halo-3-trifluoromethylbenzonitriles

AUTHOR(S): Hino, Tetsuo; Namie, Tsutomu; Nakamura, Hiroyuki; Yonezawa, Noriyuki

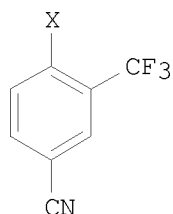
CORPORATE SOURCE: Department of Organic and Polymer Materials Chemistry, Tokyo University of Agriculture and Technology, Koganei-shi, 184-8588, Japan

SOURCE: Nippon Kagaku Kaishi (2002), (2), 219-222
CODEN: NKAKB8; ISSN: 0369-4577

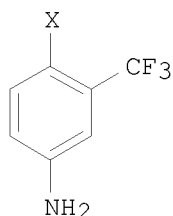
Updated Search

STNsdddaeatat

PUBLISHER: Nippon Kagakkai
DOCUMENT TYPE: Journal
LANGUAGE: Japanese
OTHER SOURCE(S): CASREACT 137:124967
GI



I



II

AB An effective synthesis of a series of fluorine-containing polyfunctional aromatic compds., 4-halo-3-trifluoromethylbenzonitriles (I; X = Cl, Br, F) from 4-halo-3-trifluoromethylanilines (II; X = same as above) was performed by the use of some tetrahedral copper-cyano complexes as Sandmeyer cyanating reagents and sublimation in the isolation and purification process. In the conversion of the corresponding diazonium salts to the target benzonitrile derivs. I, three tetrahedral tetracyanocopper complexes, K₃[Cu(CN)₄], Na₃[Cu(CN)₄], and K₂[Cu(CN)₄·NH₃], proved to be effective as Sandmeyer cyanating reagents though copper(I) cyanide gave the target compds. I in moderate yields. Furthermore, the yields of nitriles I in the reaction with the copper(I) complexes were rather higher than that with the copper(II) complex, K₂[Cu(CN)₄·NH₃]. The conversion reactions to nitriles I are suggested to proceed through an S_NAr mechanism.

OS.CITING REF COUNT: 5 THERE ARE 5 CAPLUS RECORDS THAT CITE THIS RECORD (5 CITINGS)

L13 ANSWER 5 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1998:812206 HCAPLUS

DOCUMENT NUMBER: 130:139155

TITLE: Arenediazonium tetrachlorocuprates(II). Modification of the Meerwein and Sandmeyer reactions

AUTHOR(S): Obushak, Mykola D.; Lyakhovych, Mykhaylo B.; Ganushchak, Mykola I.

CORPORATE SOURCE: Department of Organic Chemistry, Lviv State University, Lvov, 290602, Ukraine

SOURCE: Tetrahedron Letters (1998), 39(51), 9567-9570
CODEN: TELEAY; ISSN: 0040-4039

PUBLISHER: Elsevier Science Ltd.

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 130:139155

AB In the Cu-catalyzed reactions of arenediazonium chlorides with unsatd. compds., arenediazonium tetrachlorocuprates(II) are formed as intermediates. A general method of preparation of these complexed diazonium salts is described. In polar solvents these salts undergo chlorinative

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dediazonation to give chloroarenes in high yield. The reaction of an arenediazonium tetrachlorocuprate(II) with an activated alkene results in the same products as the Meerwein reaction. A radical cation mechanism for this reaction is proposed.

OS.CITING REF COUNT: 13 THERE ARE 13 CAPLUS RECORDS THAT CITE THIS RECORD (13 CITINGS)
REFERENCE COUNT: 23 THERE ARE 23 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L13 ANSWER 6 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1995:985193 HCAPLUS

DOCUMENT NUMBER: 124:175098

ORIGINAL REFERENCE NO.: 124:32463a,32466a

TITLE: Sandmeyer reactions. Part 3.
Estimation of absolute rate constants for the transfer of chloride ligands from CuII to 2-benzoylphenyl radical (Pschorr radical clock) and further investigations of the relative rates of transfer of chloride and water ligands to other substituted phenyl radicals

AUTHOR(S): Hanson, Peter; Hammond, Roger C.; Gilbert, Bruce C.; Timms, Allan W.

CORPORATE SOURCE: Dep. Chemistry, Univ. York, York, YO1 5DD, UK

SOURCE: Journal of the Chemical Society, Perkin Transactions 2: Physical Organic Chemistry (1995), (12), 2195-202
CODEN: JCPKBH; ISSN: 0300-9580

PUBLISHER: Royal Society of Chemistry

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Absolute rate consts. have been estimated for the transfer of chloride ligands from $\text{CuCl}(\text{OH}_2)_5^+$, $\text{CuCl}_2(\text{OH}_2)_4$ and $\text{CuCl}_3(\text{OH}_2)_3^-$ to 2-benzoylphenyl radical in aqueous solution at ionic strength 5 mol dm⁻³ by comparison with its rate of cyclization. The radical reacts with the monochlorocopper(II) complex with $k = (8.8 \pm 1.3) \times 10^6 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$, but it does not discriminate between the di- and tri-chloro complexes, $k = (1.0 \pm 0.15) \times 10^8 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$. Relative rate consts. for the transfer of chloride and water ligands to a series of 2-, 3- and 4-substituted Ph radicals have also been obtained; three copper complexes are implicated in chloride transfer for the conditions examined. Radicals with 3- and 4-substituents discriminate the monochlorocopper(II) complex from the two more highly chlorinated complexes, the discrimination increasing the more nucleophilic and reactive the radical. Radicals with strongly electron-withdrawing 2-substituents (CN, F) do not discriminate between any of the chlorocopper(II) complexes. A mechanistic rationale of the observations is proposed.

OS.CITING REF COUNT: 7 THERE ARE 7 CAPLUS RECORDS THAT CITE THIS RECORD (7 CITINGS)

L13 ANSWER 7 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1995:733290 HCAPLUS

DOCUMENT NUMBER: 123:111866

ORIGINAL REFERENCE NO.: 123:19988h,19989a

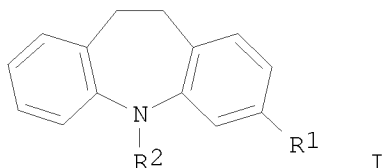
TITLE: New method for production 3-chloro-, 3-bromo-, and 3-iodo-10,11-dihydro-5H-dibenz[b,f]azepine derivatives involving one-step diazotization/Sandmeyer reaction of 5-acetyl-3-aminoiminodibenzyl

Updated Search

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INVENTOR(S): Hosztafi, Sandor; Galamb, Vilmos; Csende, Ferenc;
Nagy, Janosne; Frank, Laszlo
PATENT ASSIGNEE(S): Alkaloida Vegyeszeti Gyar Rt., Hung.
SOURCE: Hung. Teljes, 9 pp.
CODEN: HUXXB
DOCUMENT TYPE: Patent
LANGUAGE: Hungarian
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
HU 67041	A2	19950130	HU 1992-3980	19921216
PRIORITY APPLN. INFO.:			HU 1992-3980	19921216
OTHER SOURCE(S):	CASREACT 123:111866; MARPAT 123:111866			
GI				



AB 10,11-Dihydro-5H-dibenz[b,f]azepine derivs. I (R1 = Cl, Br, I; R2 = H, acyl) are prepared by one-step diazotization/Sandmeyer reaction of 5-acetyl-3-aminoiminodibenzyl (I; R1 = NH2, R2 = Ac) with isopentyl nitrite and anhydrous CuCl2, CuBr2, or CuI. Thus, e.g., anhydrous CuCl2 was added to isopentyl nitrite in MeCN; then 3-amino-5-acetylaminodibenzyl was added so that the temperature did not exceed 30°; mixing for 2 h followed by workup afforded 70% 5-acetyl-3-chloroiminodibenzyl.

OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD
(1 CITINGS)

L13 ANSWER 8 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1991:558229 HCAPLUS
DOCUMENT NUMBER: 115:158229
ORIGINAL REFERENCE NO.: 115:27079a,27082a
TITLE: Sandmeyer reactions. Part 1. A comparative study of the transfer of halide and water ligands from complexes of copper(II) to aryl radicals
AUTHOR(S): Hanson, Peter; Jones, Jason R.; Gilbert, Bruce C.; Timms, Allan W.
CORPORATE SOURCE: Dep. Chem., Univ. York, Heslington/York, YO1 5DD, UK
SOURCE: Journal of the Chemical Society, Perkin Transactions 2: Physical Organic Chemistry (1972-1999) (1991), (7), 1009-17
CODEN: JCPKBH; ISSN: 0300-9580
DOCUMENT TYPE: Journal

Updated Search

STNsdddaeatat

LANGUAGE: English

AB Evidence was presented indicating that the homolytic formation of phenols under Sandmeyer conditions was directly comparable with the well-known halogenation reactions. Sandmeyer hydroxylation and halogenation are competitive processes; however, transfer of a halide ligand occurs more rapidly than that of a water ligand, and at high halide concns. phenol formation is minor, consistent with long known synthetic results. Relative rate data for ligand transfers to 4-substituted Ph radicals from complexes were given. LFER were examined using σ I and σ R.

OS.CITING REF COUNT: 12 THERE ARE 12 CAPLUS RECORDS THAT CITE THIS RECORD (12 CITINGS)

L13 ANSWER 9 OF 9 HCAPLUS COPYRIGHT 2010 ACS on STN

ACCESSION NUMBER: 1977:452884 HCAPLUS

DOCUMENT NUMBER: 87:52884

ORIGINAL REFERENCE NO.: 87:8367a,8370a

TITLE: Alkyl nitrite-metal halide deamination reactions. 2. Substitutive deamination of arylamines by alkyl nitrites and copper(II) halides. A direct and remarkably efficient conversion of arylamines to aryl halides

AUTHOR(S): Doyle, Michael P.; Siegfried, Bernard; Dellaria, Joseph F., Jr.

CORPORATE SOURCE: Dep. Chem., Hope Coll., Holland, MI, USA

SOURCE: Journal of Organic Chemistry (1977), 42(14), 2426-31
CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal

LANGUAGE: English

OTHER SOURCE(S): CASREACT 87:52884

AB Arylamines (15) with 1 molar equivalent tert-BuONO and 0.5 molar equivalent CuCl₂

or CuBr₂ resulted in substitutive deamination to give the aryl chlorides or bromides. E.g., O₂NC₆H₄NH₂ and 1-aminonaphthalene with CuCl₂ and tert-BuONO in MeCN gave 99.5% p-O₂NC₆H₄Cl and 82% 1-chloronaphthalene, resp. In the reactions using CuBr₂, bromination at the o- or p-positions also occurred. The deamination-substitution procedure using Cu(II) halides was compared with that of the Sandmeyer reaction which uses Cu(I) halides.

OS.CITING REF COUNT: 102 THERE ARE 102 CAPLUS RECORDS THAT CITE THIS RECORD (103 CITINGS)

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